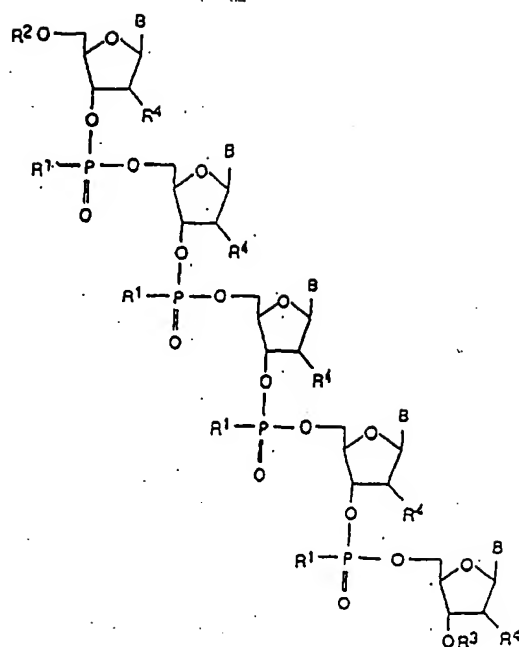


This listing of claims will replace all prior versions, and listings, of claims in the application.

In the Claims:

1. (CURRENTLY AMENDED) An antisense oligonucleotide selected from the group consisting of ~~[[]]~~ the sequence 5'- TTG CAT AAA CCC AAG GAG - 3' (SEQ ID NO: 1) and modifications thereof, and a ~~[[]]~~ fragment having at least 8 nucleotides of the sequence 5'- TTG CAT AAA CCC AAG GAG - 3' (SEQ ID NO: 1) and modifications thereof.
2. (CURRENTLY AMENDED) The antisense-oligonucleotide according to claim 1 wherein the modification ~~concerns one or more of the sugar moieties, the bases and/or the internucleotide linkages and/or~~ comprises a modified sugar moiety, a modified base, a modified internucleotide linkage, and/or coupling the oligonucleotide to an enhancer of uptake and/or inhibitory activity, and combinations thereof.
3. (ORIGINAL) The antisense-oligonucleotide according to claim 2 wherein the antisense-oligonucleotide is a phosphorothioate oligodeoxynucleotide.
4. (CURRENTLY AMENDED) The antisense-oligonucleotides according to claim 1 with the ~~respective~~ structure:



wherein

[[-]] B = the bases A, C, G or T in oligodeoxy-ribonucleotides or ~~accordingly~~ the bases A, C, G or U in oligo-ribonucleotides;

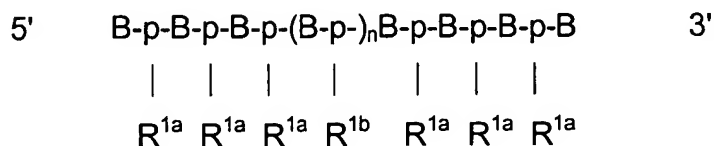
[[-]] $R^1 = O^-M^+$ ($M^+ = Na^+$ or H^+), S^-M^+ ($M^+ = Na^+$ or H^+), CH_3 , C_2H_5 , OCH_3 , or OC_2H_5 ;

[[-]] R^2 and/or R^3 are covalently coupled cholesterol, poly(L)lysine, transferrin or H ;

[[-]] $R^4 = H$, F , CH_3 , C_2H_5 , OH , OCH_3 , or OC_2H_5 ;

~~and wherein the structure is to be understood as a detail out a representation of a~~
longer nucleotide chain.

5. (CURRENTLY AMENDED) The aAntisense oligonucleotides according to claim 1 with the formula



wherein

B = the bases A, C, G or T in oligodeoxy-ribonucleotides, or ~~accordingly~~ the bases A, C, G or U in oligo-ribonucleotides;

p = internucleotide phosphate;

(B-p)_n = an oligodeoxy-ribonucleotide or oligo-ribonucleotide stretch wherein

n = 1 – 12, ~~preferably 1–11~~;

and wherein R¹, ~~referred to as encompassing~~ R^{1a} or R^{1b}, is varied at the internucleotide phosphates within one oligonucleotide[[:]] wherein

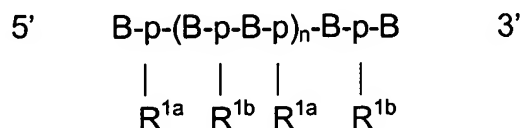
R^{1a} = S⁻M⁺, wherein all M⁺ is Na⁺ or H⁺ and R^{1b} = O⁻M⁺, wherein all M⁺ is Na⁺ or H⁺; or

R^{1a} = CH₃ and R^{1b} = O⁻M⁺, wherein all M⁺ is Na⁺ or H⁺; or

R^{1a} = S⁻M⁺, wherein all M⁺ is Na⁺ or H⁺S and R^{1b} = CH₃; or

R^{1a} = CH₃ and R^{1b} = S⁻M⁺, wherein all M⁺ is Na⁺ or H⁺.

6. (CURRENTLY AMENDED) The aAntisense oligonucleotides according to claim 1 with the formula



wherein

B = one of the bases A, C, G or T ~~comprised~~ in oligodeoxy-ribonucleotides or ~~accord-~~
ingly one of the bases A, C, G or U ~~comprised~~ in oligo-ribonucleotides depending on a
gene sequence;

p = internucleotide phosphate;

(B-p-B-p)_n = an oligodeoxy-ribonucleotide or oligo-ribonucleotide stretch wherein n = 2
– 8, preferably 3–7;

and wherein R¹ is alternated at the internucleotide phosphates within one oligonucleo-
tide[[:]] wherein

R^{1a} = S⁺M⁺, wherein all M⁺ is Na⁺ or H⁺ and R^{1b} = O⁻M⁺, wherein all M⁺ is Na⁺ or H⁺; or

R^{1a} = CH₃ and R^{1b} = O⁻M⁺, wherein all M⁺ is Na⁺ or H⁺; or

R^{1a} = S⁺M⁺, wherein all M⁺ is Na⁺ or H⁺ and R^{1b} = CH₃.

7. (CURRENTLY AMENDED) Use of the antisense-oligonucleotides according to claim 1
for at least one of the inhibition of expression and/or functional activity of melanoma
inhibitory activity (MIA), ~~and/or~~ reducing invasion and/or metastasis, ~~and/or~~ or stimu-
lating immune cells and/or the immune system.
8. (ORIGINAL) A pharmaceutical composition comprising an antisense-oligonucleotide
according to claim 1.
9. (CURRENTLY AMENDED) The pharmaceutical composition according to claim 8
wherein the antisense-oligonucleotide is integrated into a DNA delivery system com-
prising viral and/or non-viral vectors together with lipid acids or derivatives thereof
selected from the group consisting of anionic lipids, cationic lipids, non-cationic lip-
ids, and mixtures thereof.

10. (CURRENTLY AMENDED) The pharmaceutical composition according to claim 8 further comprising ~~additionally~~ an immunostimulatory agent.
11. (CURRENTLY AMENDED) The pharmaceutical composition according to claim 10 wherein the ~~additionally~~ immunostimulatory agent is selected from the group consisting of cytokines, inhibitors of expression and/or function of interleukin-10, inhibitors of expression and/or function of transforming growth factor beta (TGF- β), inhibitors of expression and/or function of Prostaglandin B2, inhibitors of expression and/or function of receptors for Prostaglandin E2, ~~and/or~~ inhibitors of VEGF, and combinations thereof.
12. (CURRENTLY AMENDED) The use of the pharmaceutical composition according to ~~one of the claims 8-14~~ claim 8 in a method for the ~~preparation of a medicament~~ prevention and/or the treatment of at least one of neoplasms, infections, or immunosuppressive disorders.
13. (CURRENTLY AMENDED) The use of the pharmaceutical composition according to ~~one of the claims 8-14~~ claim 8 in a method for the ~~preparation of a medicament~~ prevention and/or the treatment of at least one disorder[[s]], neoplasm[[s]], infection[[s]], ~~and/or or~~ immunosuppressive disorder[[s]] ~~where an~~ wherein abnormal expression of MIA plays a role in the disorder, neoplasm, infection, or immunosuppressive disorder.
14. (CURRENTLY AMENDED) The use of the pharmaceutical composition according to ~~the claims 8-14~~ claim 8 in a method for the ~~preparation of a medicament~~ prevention

[[or]] and/or the treatment of neoplasms and/or disorders selected from the group consisting of melanoma, gastrointestinal carcinoma, breast cancer, pancreatic cancer, [[ovarial]] ovarian carcinoma, chondrosarcoma, spinal diseases, cervical myelopathy, lumbar canal stenosis, lumbar disc herniation, rheumatoid arthritis, osteoarthritis, HLA-27-associated oligoarthritis, psoriatic arthritis, [[and]] rheumatic arthritis, cartilage damage, [[or]] joint destruction, and combinations thereof.

15. (CURRENTLY AMENDED) A diagnostic composition comprising an antisense-oligonucleotide according to ~~one of the claims 1-7~~ either claim 1 or claim 2.
16. (NEW) The composition of claim 5 wherein (B-p-)_n = an oligodeoxy-ribonucleotide or oligo-ribonucleotide stretch wherein n= 1 – 11.
17. (NEW) The composition of claim 6 wherein (B-p-B-p)_n = an oligodeoxy-ribonucleotide or oligo-ribonucleotide stretch wherein n= 3 – 7.